## IN THE CLAIMS

- 1-17. (Cancelled)
- 18. (Currently amended) A method of detecting measuring activity of an NAD+ utilizing enzyme, comprising:

incubating the enzyme with NAD+ and a substrate for the enzyme; quantifying any remaining NAD+ by the method of claim 12 converting any remaining NAD+ to a fluorescent compound; and measuring an amount of fluorescence of the fluorescent compound.

19. (Original) The method of claim 18, wherein the fluorescent compound is compound 1:

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20. (Original) The method of claim 18, wherein the converting comprises: mixing NAD+ with acetophenone and base, to form a mixture; and reacting the mixture with acid.

- 21. (Original) The method of claim 20, wherein the base is a solution of KOH.
- 22. (Original) The method of claim 20, wherein the acid comprises formic acid.
- 23. (Original) The method of claim 20, wherein the fluorescent compound is compound 1:

- 24. (Original) The method of claim 18, wherein the enzyme is PARP.
- 25. (Currently amended) A method of determining whether a compound is an inhibitor of an NAD+ utilizing enzyme, comprising:

comparing an amount of NAD+ consumed during reaction of the enzyme with a substrate for the enzyme measuring activity of the enzyme by the method of claim 18, with and without the compound; and

comparing the measured activity of the enzyme with the compound and the measured activity of the enzyme without the compound

wherein the amount of NAD+ not consumed is measured by the method of claim 12.

(Original) The method of claim 25, wherein the fluorescent compound is 26. compound 1:

- 27. (Original) The method of claim 25, wherein the converting comprises: mixing NAD+ with acetophenone and base, to form a mixture; and reacting the mixture with acid.
- 28. (Original) The method of claim 27, wherein the base is a solution of KOH.
- 29. (Original) The method of claim 27, wherein the acid comprises formic acid.

30. (Original) The method of claim 27, wherein the fluorescent compound is compound 1:

- 31. (Original) The method of claim 25, wherein the enzyme is PARP.
- 32. (Original) The method of claim 27, wherein the enzyme is PARP.
- 33. (Currently amended) A method of detecting a genetic deficiency in an NAD+ utilizing enzyme in a patient, comprising:

measuring activity of the enzyme from the patient and a control enzyme, by the method of claim 18; and

comparing an amount of NAD+ consumed during reaction of an the measured activity of the enzyme from the patient with a substrate for the enzyme, with an amount of NAD+ consumed during reaction of a and the measured activity of the control enzyme with the substrate;

wherein the amount of NAD+ not consumed is measured by the method of claim 12.

34. (Original) The method of claim 33, wherein the fluorescent compound is compound 1:

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- 35. (Original) The method of claim 33, wherein the converting comprises: mixing NAD+ with acetophenone and base, to form a mixture; and reacting the mixture with acid.
- 36. (Original) The method of claim 35, wherein the base is a solution of KOH.
- 37. (Original) The method of claim 35, wherein the acid comprises formic acid.

(Original) The method of claim 35, wherein the fluorescent compound is 38. compound 1:

39. (Original) The method of claim 33, wherein the NAD+ utilizing enzyme is long-chain 3-hydroxyacyl-CoA dehydrogenase.

40-53. (Cancelled)